

~~2-Aminoaryloxazole compounds for the treatment of diseases~~

5 The present invention relates to novel compounds selected from 2-aminoaryloxazoles that selectively modulate, regulate, and/or inhibit signal transduction mediated by certain native and/or mutant tyrosine kinases implicated in a variety of human and animal diseases such as cell proliferative, metabolic, allergic, and degenerative disorders. More particularly, these compounds are potent and selective c-kit, bcr-abl,
10 FGFR3 and/or Flt-3 inhibitors.

Tyrosine kinases are receptor type or non-receptor type proteins, which transfer the terminal phosphate of ATP to tyrosine residues of proteins thereby activating or inactivating signal transduction pathways. These proteins are known to be involved in
15 many cellular mechanisms, which in case of disruption, lead to disorders such as abnormal cell proliferation and migration as well as inflammation.

As of today, there are about 58 known receptor tyrosine kinases. Included are the well-known VEGF receptors (Kim et al., Nature 362, pp. 841-844, 1993), PDGF
20 receptors, c-kit, Flt-3 and the FLK family. These receptors can transmit signals to other tyrosine kinases including Src, Raf, Frk, Btk, Csk, Abl, Fes/Fps, Fak, Jak, Ack, etc.

Among tyrosine kinase receptors, c-kit is of special interest. Indeed, c-kit is a key
25 receptor activating mast cells, which have proved to be directly or indirectly implicated in numerous pathologies for which the Applicant filed WO 03/004007, WO 03/004006, WO 03/003006, WO 03/003004, WO 03/002114, WO 03/002109, WO 03/002108, WO 03/002107, WO 03/002106, WO 03/002105, WO 03/039550, WO 03/035050, WO 03/035049, WO 03/0720090, WO 03/072106 and
30 IB2004/000907, as well as US 60/495,088.

It was found that mast cells present in tissues of patients are implicated in or contribute to the genesis of diseases such as autoimmune diseases (rheumatoid arthritis, inflammatory bowel diseases (IBD)) allergic diseases, bone loss, cancers such as solid tumors, leukaemia and GIST, tumor angiogenesis, inflammatory diseases, interstitial cystitis, mastocytosis, graft-versus-host diseases, infection diseases, metabolic disorders, fibrosis, diabetes and CNS diseases. In these diseases, it has been shown that mast cells participate in the destruction of tissues by releasing a cocktail of different proteases and mediators such as histamine, neutral proteases, lipid-derived mediators (prostaglandins, thromboxanes and leucotrienes), and various cytokines (IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-8, TNF- α , GM-CSF, MIP-1a, MIP-1b, MIP-2 and IFN- γ).

The c-kit receptor can also be constitutively activated by mutations leading to abnormal cell proliferation and development of diseases such as mastocytosis (D816V mutation) and various cancers such as GIST (c-kit Δ 27, a juxtamembrane deletion).

Furthermore, 60% to 70% of patients presenting with AML have blasts which express c-kit, the receptor for stem cell factor (SCF) (Broudy, 1997). SCF promotes growth of hematopoietic progenitors, and act as a survival factor for AML blasts. In some cases (1 to 2%) of AML, a mutation in a conserved residue of the kinase domain (Kit816) resulting in constitutive activation of c-kit has been described (Beghini et al., 2000; Longley et al., 2001). This gain of function mutation (Asp to Val/Tyr substitution) has been identified in mast cell leukemic cell lines and in samples derived from patients with mastocytosis (Longley et al., 1996).

We have studied about 300 patients afflicted with systemic mastocytosis and we have shown that the Kit816 mutation is expressed in about 60% of cases. In this regard, we filed IB 2004/000907 which relates to tailored treatment of the different forms of mastocytosis depending on the presence or absence of the Kit816 mutation.

In view of the above, we have proposed to target c-kit to deplete the mast cells responsible for these disorders.

In addition, 60 to 80% of AML blasts express a similar receptor Flt3, the receptor for
5 Flt3 ligand and in a high percentage of ALL. Both ligand and receptor have been
identified by Hannum et al., 1994 and Rosnet et al., 1991. Like the c-Kit, Flt3
mediates differentiation and proliferation of normal hematopoietic stem cells and
mediates proliferation and survival signals in AML blasts. Although Flt3 is most
commonly expressed in the wild type form, the leukemic clone of 30 to 35% of
10 patients with AML (Nakao et al., 1996), expresses a mutated form of Flt3 that
contains an Internal Tandem Duplication (Flt3ITD) of the juxtamembrane domain
coding sequence. This mutation leads to constitutive activation of the receptor and
autonomous cytokine-independent growth. It has also been reported that a cohort of
AML patients (~7%) contains mutations in the activation loop of Flt3 at amino acid
15 position Asp835 (FLT3835) (Yamamoto et al., 2001). This mutation occurs at the
corresponding position in c-kit (Kit816) described above, leading kinases to adopt an
activated configuration. Flt3 mutations have also been reported at a frequency of 15%
in secondary AML and may be associated with disease progression or relapse of
AML.

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Here, we provide for the first time inhibitors of Flt3ITD for treating for example 30 to
35% of patients with AML presenting this mutation.

Among our compounds, we also have found inhibitors of FGFR3 which is
25 responsible for several lethal cancers.

Many different compounds have been described as tyrosine kinase inhibitors, for
example, bis monocyclic, bicyclic or heterocyclic aryl compounds (WO 92/20642),
vinylene-azaindole derivatives (WO 94/14808), 1-cyclopropyl-4-pyridyl-quinolones

(US 5,330,992), styryl compounds (US 5,217,999), styryl-substituted pyridyl compounds (US 5,302,606), selenoindoles and selenides (WO 94/03427), tricyclic polyhydroxylic compounds (WO 92/21660), benzylphosphonic acid compounds (WO 91/15495), pyrimidine derivatives (US 5,521,184 and WO 99/03854), indolinone derivatives and pyrrole-substituted indolinones (US 5,792,783, EP 934 931, US 5,834,504, US 5,883,116, US 5,883,113, US 5, 886,020, WO 96/40116 and WO 00/38519), as well as bis monocyclic, bicyclic aryl and heteroaryl compounds (EP 584 222, US 5,656,643 and WO 92/20642), quinazoline derivatives (EP 602 851, EP 520 722, US 3,772,295 and US 4,343,940) and aryl and heteroaryl quinazoline (US 5,721,237, US 5,714,493, US 5,710,158 and WO 95/15758).

There are hundreds of tyrosine kinases in mammalian cells that are more or less prone to be modulated by the compounds cited above. The problem is that a tyrosine kinase inhibitor has to be very specific to one or very few kinases to avoid toxicity and side effects on the long run. None of these prior art tyrosine kinase inhibitors provides a solution for this problem. Besides, none of these compounds have been described as potent and selective inhibitors of c-kit or of the c-kit pathway, nor highly specific bcr-abl, FGFR and/or Flt-3 inhibitors.

The present invention provides potent and selective compounds capable of inhibiting wild type and/or mutated c-kit, as well as subsets of compounds inhibiting c-kit, bcr-abl, FGFR3 and/or Flt-3.

In connection with the present invention, we have found that compounds corresponding to the 2-aminoaryloxazoles are potent and selective inhibitors of c-kit, bcr-abl, FGFR3 and/or Flt-3. These compounds are good candidates for treating diseases such as autoimmune diseases, inflammatory diseases, cancers and mastocytosis. Compounds of the invention displaying inhibitory activity on Flt3 are particularly suitable for treating different forms of leukemia, such as AML. Compounds of the invention displaying inhibitory activity on FGFR3 constitute a

breakthrough for treating lethal cancers such as bladder cancer, myeloma 414 and ariways cancers.

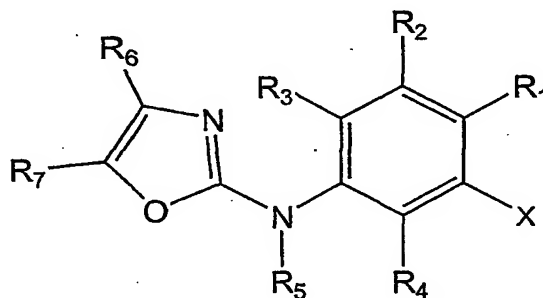
Description

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Therefore, the present invention relates to compounds belonging to the 2-aminoaryloxazoles. These compounds are capable of selectively inhibiting signal transduction involving the tyrosine phosphokinase c-kit, bcr-abl, Flt-3 and mutant
10 forms thereof.

In a first embodiment, the invention is aimed at compounds of formula I, which may represent either free base forms of the substances or pharmaceutically acceptable salts thereof:

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FORMULA I

20 Substituents R₁ - R₇ and X in Formula I are defined as follows:

R₁, R₂, R₃ and R₄ each independently are selected from hydrogen, halogen (selected from F, Cl, Br or I), a linear or branched alkyl group containing from 1 to 10 carbon atoms and optionally substituted with one or more heteroatoms such as halogen
25 (selected from F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant basic nitrogen functionality, as well as trifluoromethyl, C₁₋₆alkyloxy, amino, C₁₋₆alkylamino, di(C₁₋₆alkyl)amino, carboxyl, cyano, nitro, formyl, hydroxy, and CO-

6.

R, COO-R, CONH-R, SO₂-R, and SO₂NH-R wherein R is a linear or branched alkyl group containing from 1 to 10 carbon atoms and optionally substituted with at least one heteroatom, notably a halogen (selected from F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant basic nitrogen functionality.

5 R₅ is one of the following:

(i) hydrogen, or

(ii) a linear or branched alkyl group containing from 1 to 10 carbon atoms and optionally substituted with one or more heteroatoms such as halogen (selected from F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant
10 basic nitrogen functionality, or

(iii) CO-R₈ or COOR₈ or CONHR₈ or SO₂R₈ wherein R₈ may be

- a linear or branched alkyl group containing from 1 to 10 carbon atoms and optionally substituted with one or more heteroatoms such as halogen (selected from F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant
15 basic nitrogen functionality, or

- an aryl group such as phenyl or a substituted variant thereof bearing any combination, at any one ring position, of one or more substituents such as halogen (selected from F, Cl, Br or I), alkyl groups containing from 1 to 10 carbon atoms and optionally substituted with one or more heteroatoms such as halogen (selected from
20 F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant basic nitrogen functionality; as well as trifluoromethyl, C₁₋₆alkyloxy, carboxyl, cyano, nitro, formyl, hydroxy, C₁₋₆alkylamino, di(C₁₋₆alkyl)amino, and amino, the latter nitrogen substituents optionally in the form of a pendant basic nitrogen functionality; as well as CO-R, COO-R, CONH-R, SO₂-R, and SO₂NH-R wherein R is a linear or
25 branched alkyl group containing from 1 to 10 carbon atoms and optionally substituted with at least one heteroatom, notably a halogen (selected from F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant basic nitrogen functionality, or

- a heteroaryl group such as a pyridyl, pyrimidinyl, pyrazinyl, pyridazinyl, thienyl, thiazolyl, imidazolyl, pyrazolyl, pyrrolyl, furanyl, oxazolyl, isoxazolyl, triazolyl, tetrazolyl, indolyl, benzimidazole, quinolinyl group, which may additionally
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bear any combination, at any one ring position, of one or more substituents such as halogen (selected from F, Cl, Br or I), alkyl groups containing from 1 to 10 carbon atoms and optionally substituted with one or more heteroatoms such as halogen (selected from F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant basic nitrogen functionality; as well as trifluoromethyl, C₁₋₆alkoxy, carboxyl, cyano, nitro, formyl, hydroxy, C₁₋₆alkylamino, di(C₁₋₆alkyl)amino, and amino, the latter nitrogen substituents optionally in the form of a basic nitrogen functionality; as well as CO-R, COO-R, CONH-R, SO₂-R, and SO₂NH-R wherein R is a linear or branched alkyl group containing from 1 to 10 carbon atoms and optionally substituted with at least one heteroatom, notably a halogen (selected from F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant basic nitrogen functionality.

R6 and R7 each independently are selected from:

- i) hydrogen, a halogen (selected from F, Cl, Br or I), or
- ii) an alkyl¹ group defined as a linear, branched or cycloalkyl group containing from 1 to 10 carbon atoms, or from 2 or 3 to 10 carbon atoms, (for example methyl, ethyl, propyl, butyl, pentyl, hexyl...) and optionally substituted with one or more heteroatoms such as halogen (selected from F, Cl, Br or I), oxygen, and nitrogen (the latter optionally in the form of a pendant basic nitrogen functionality); as well as trifluoromethyl, carboxyl, cyano, nitro, formyl; as well as CO-R, COO-R, CONH-R, SO₂-R, and SO₂NH-R wherein R is a linear or branched alkyl group containing 1 to 10 carbon atoms, or from 2 or 3 to 10 carbon atoms, (for example methyl, ethyl, propyl, butyl, pentyl, hexyl...) and optionally substituted with at least one heteroatom, notably a halogen (selected from F, Cl, Br or I), oxygen, and nitrogen, the latter optionally in the form of a pendant basic nitrogen functionality ; as well as a cycloalkyl or aryl¹ or heteroaryl¹ group optionally substituted by a pendant basic nitrogen functionality,

or

(iii) an aryl¹ group defined as phenyl or a substituted variant thereof bearing any combination, at any one ring position, of one or more substituents such as

- halogen(selected from I, F, Cl or Br);
- an alkyl¹ group;
- 5 - a cycloalkyl, aryl or heteroaryl group optionally substituted by a pendant basic nitrogen functionality;
- trifluoromethyl, O-alkyl¹, carboxyl, cyano, nitro, formyl, hydroxy, NH-alkyl¹, N(alkyl¹)(alkyl¹), and amino, the latter nitrogen substituents optionally in the form of a basic nitrogen functionality;
- 10 - NHCO-R or NHCOO-R or NHCONH-R or NHSO₂-R or NHSO₂NH-R or CO-R or COO-R or CONH-R or SO₂-R or SO₂NH-R wherein R corresponds to hydrogen, alkyl¹, aryl or heteroaryl, or

(iv) a heteroaryl¹ group defined as a pyridyl, pyrimidinyl, pyrazinyl, pyridazinyl, thienyl, thiazolyl, imidazolyl, pyrazolyl, pyrrolyl, furanyl, oxazolyl, isoxazolyl , triazolyl, tetrazolyl, indolyl, benzimidazole, quinolinyl group, which may additionally bear any combination, at any one ring position, of one or more substituents such as

- halogen (selected from F, Cl, Br or I);
- an alkyl¹ group;
- 20 - a cycloalkyl, aryl or heteroaryl group optionally substituted by a pendant basic nitrogen functionality,
- trifluoromethyl, O-alkyl¹, carboxyl, cyano, nitro, formyl, hydroxy, NH-alkyl¹, N(alkyl¹)(alkyl¹), and amino, the latter nitrogen substituents optionally in the form of a basic nitrogen functionality;
- 25 - NHCO-R or NHCOO-R or NHCONH-R or NHSO₂-R or NHSO₂NH-R or CO-R or COO-R or CONH-R or SO₂-R or SO₂NH-R wherein R corresponds to hydrogen, alkyl¹, or

(v) an O-aryl¹, or NH-aryl¹, or O-heteroaryl¹ or NH-heteroaryl¹ group

(vi) trifluoromethyl, O-alkyl¹, carboxyl, cyano, nitro, formyl, hydroxy, NH-alkyl¹, N(alkyl¹)(alkyl¹), and amino, the latter nitrogen substituents optionally in the form of a basic nitrogen functionality, or

(vi) NHCO-R or NHCOO-R or NHCONH-R or NHSO₂-R or NHSO₂NH-R or CO-R
 5 or COO-R or CONH-R or SO₂-R or SO₂NH-R wherein R corresponds to hydrogen, alkyl¹, aryl or heteroaryl.

Substituent X is:

-NR₉R₁₀, wherein R₉ and / or R₁₀ are hydrogen or:

- 10 i) an alkyl¹ group, CF₃ or
 ii) an aryl¹, heteroaryl¹ or cycloalkyl group optionally substituted by a pendant basic nitrogen functionality, or
 iii) a CO-R, COO-R, CON-RR' or SO₂-R, where R and R' are a hydrogen, alkyl¹,
 aryl¹ or heteroaryl¹, optionally substituted by a pendant basic nitrogen functionality;

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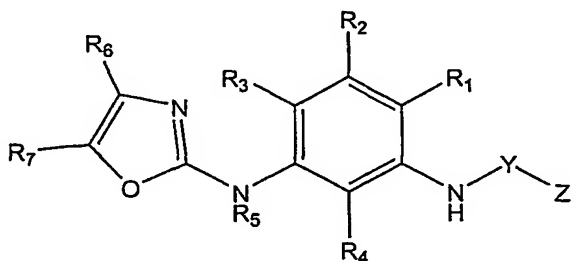
or:

-CO-NR₉R₁₀, wherein R₉ and / or R₁₀ are hydrogen or:

- i) an alkyl¹ group, CF₃ or
 ii) an aryl¹, heteroaryl¹ or cycloalkyl group optionally substituted by a pendant basic
 20 nitrogen functionality, or

X may also be Alkyl¹.

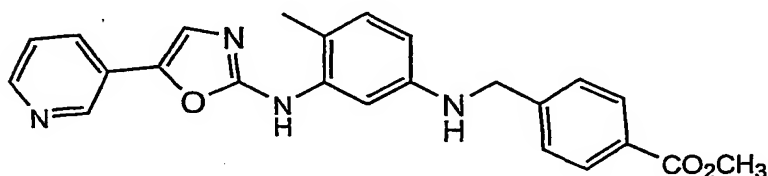
Among the particular compounds of formula I, the invention is directed to oxazol-2-
 25 yl-benzene-1,3-diamine compounds of the following formula I-2:



wherein R5 = H, Y and Z represents an hydrogen, an aryl¹ or a heteroaryl¹ group, optionally substituted by a pendant basic nitrogen functionality. R1, R2, R3, R4, R6, and R7 have the meaning as depicted above.

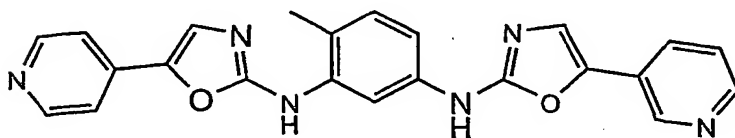
An example of preferred compounds of the above formula is depicted below:

001 : 4- {[4-Methyl-3-(4-pyridin-3-yl-oxazol-2-ylamino)-phenylamino]-methyl}-benzoic acid methyl ester



¹H NMR (CDCl₃, 300 MHz) δ = 2.15 (s, 3H) ; 3.81 (s, 3H) ; 4.35 (s, 2H) ; 6.19 (d, J = 6.0 Hz, 1H) ; 6.70 (br s, 1H) ; 6.90 (d, J = 6.0 Hz, 1H) ; 7.23 (m, 1H) ; 7.38 (d, J = 9.0, 2H) ; 7.41 (br s, 1H) ; 7.93 (d, J = 9.0, 2H) 8.88 (br s, 1H) ; 8.42 (br s, 1H) ; 8.70 (br s, 1H).

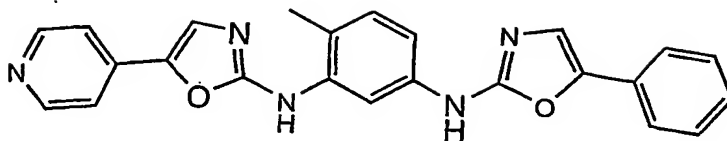
014: 4-Methyl-N1-(5-pyridin-3-yl-oxazol-2-yl)-N3-(5-pyridin-4-yl-oxazol-2-yl)-benzene-1,3-diamine



m.p. > 265°C

015: 4-Methyl-N1-(5-phenyl-oxazol-2-yl)-N3-(5-pyridin-4-yl-oxazol-2-yl)-benzene-1,3-diamine

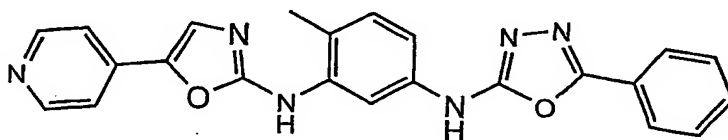
11



m.p. = 161°C

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016: 4-Methyl-N1-(5-phenyl-[1,3,4]oxadiazol-2-yl)-N3-(5-pyridin-4-yl-oxazol-2-yl)-benzene-1,3-diamine

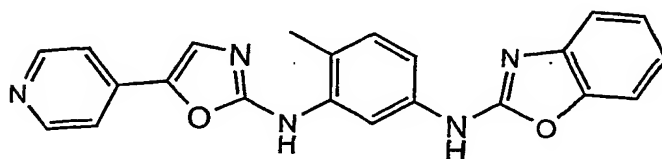


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m.p. > 265°C

017: N1-Benzooxazol-2-yl-4-methyl-N3-(5-pyridin-4-yl-oxazol-2-yl)-benzene-1,3-diamine

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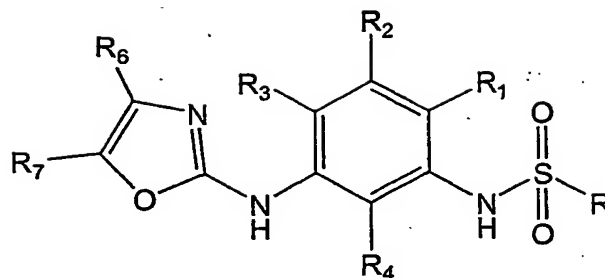


m.p. = 235°C

Among the compounds of formula I, the invention is particularly embodied by the compounds wherein R⁵ = H, X is NHSO₂R group, R is independently alkyl¹, aryl¹ or heteroaryl¹, corresponding to the family [3-(Oxazol-2-ylamino)-phenyl]-sulfonamide and the following formula I-3.

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FORMULA I-3

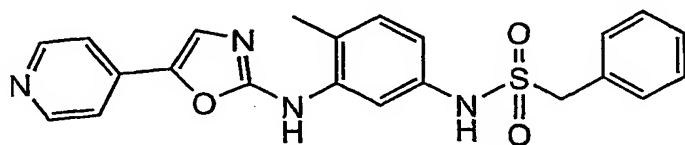
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wherein R1, R2, R3, R4, R6 and R7 have the meaning as defined above in formula I.

Examples

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057: *N*-[4-Methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-*C*-phenyl-methanesulfon -amide

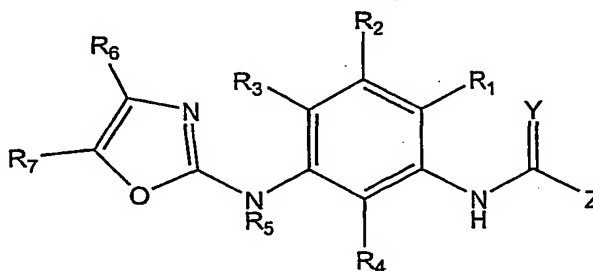


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m.p. = 190-192°C

Among the compounds of formula I, the invention is particularly embodied by the compounds of the following formula II :

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FORMULA II

Wherein R5 = H, Y is selected from O, S and Z corresponds to H, alkyl¹, or NRR', wherein R and R' are independently chosen from H or alkyl¹ or aryl¹ or heteroaryl¹, optionally substituted by a pendant basic nitrogen functionality. R1, R2, R3, R4, R6, and R7 have the meaning described above for formula I.

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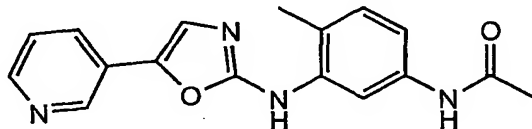
It also relates to compounds of formula II, wherein Y is selected from O, S and Z corresponds to H, NRaRb, alkyl¹, aryl¹, O-alkyl¹, or O-aryl¹ wherein Ra and Rb are independently chosen from H or alkyl¹ or aryl¹ or heteroaryl¹, optionally substituted by a pendant basic nitrogen functionality. R1, R2, R3, R4, R5, R6, and R7 have the

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Examples :

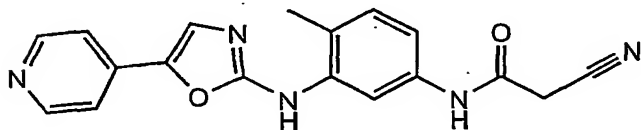
002: *N*-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-acetamide

15



m.p. = 240-242°C

20 018: 2-Cyano-*N*-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-acetamide

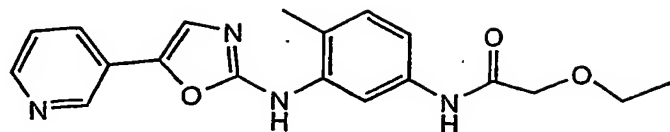


m.p. = 163°C

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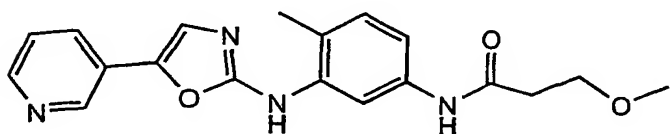
019: 2-Ethoxy-*N*-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-acetamide

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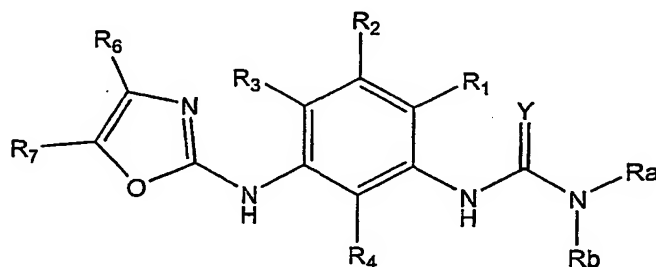
m.p. = 246-250°C

- 5 020: 3-Methoxy-N-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-propionamide



10 m.p. = 175-177°C

- Among the compounds of formula II, the invention is particularly embodied by the compounds wherein R₅ = H, Y = O or S, Z is a NR_aR_b group, corresponding to the
- 15 [3-(oxazol-2-ylamino)-phenyl]-urea or the [3-(oxazol-2-ylamino)-phenyl]-thiourea family and the following formula II-1 :



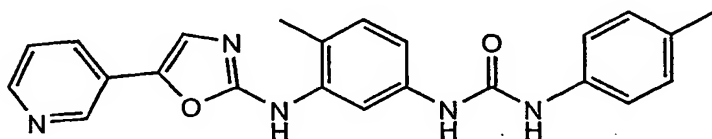
FORMULA II-1

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wherein Ra, Rb are independently chosen from H or alkyl¹ or aryl¹ or heteroaryl¹, optionally substituted by a pendant basic nitrogen functionality. R1, R2, R3, R4, R6, and R7 have the meaning described above.

5 Examples:

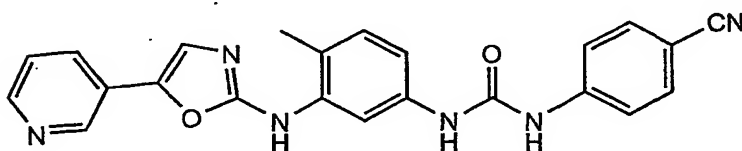
003: 1-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-*p*-tolyl-urea



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m.p. = 214°C

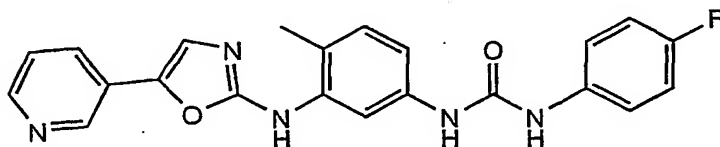
004: 1-(4-Cyano-phenyl)-3-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-urea



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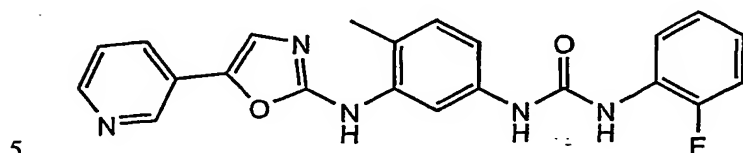
m.p. = 266°C

20 005: 1-(4-Fluoro-phenyl)-3-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-urea



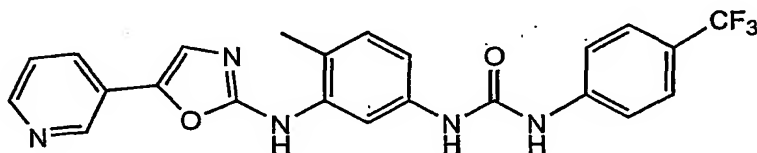
25 m.p. = 240°C

006: 1-(2-Fluoro-phenyl)-3-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-
urea



m.p. = 227°C

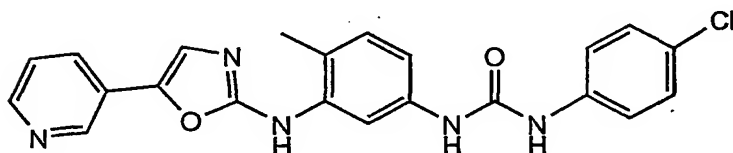
007: 1-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-(4-trifluoromethyl-
10 phenyl)-urea



m.p. = 241°C

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008: : 1-(4-Chloro-phenyl)-3-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-
urea

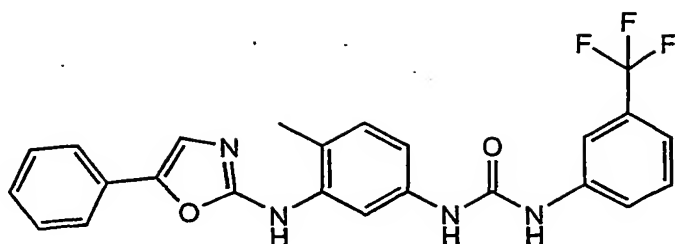


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m.p. = 220°C

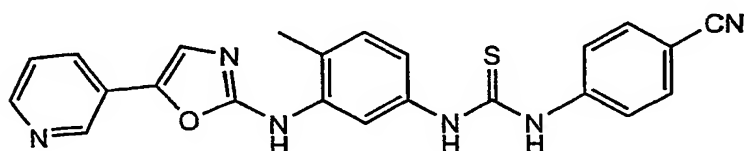
021: 1-[4-Methyl-3-(5-phenyl-oxazol-2-ylamino)-phenyl]-3-(3-trifluoromethyl-
phenyl)-urea

17



m.p. = 243°C

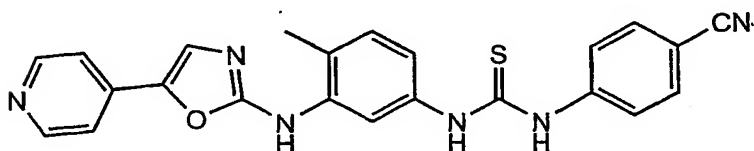
- 5 022: 1-(4-Cyano-phenyl)-3-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-thiourea



m.p. = 205°C

10

- 023: 1-(4-Cyano-phenyl)-3-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-thiourea

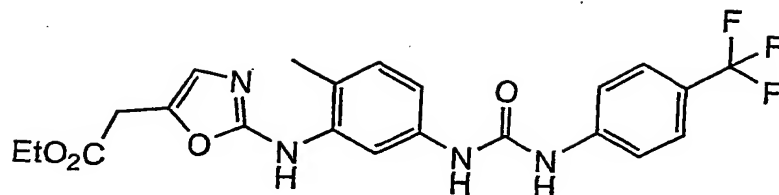


15

m.p. = 169-171°C

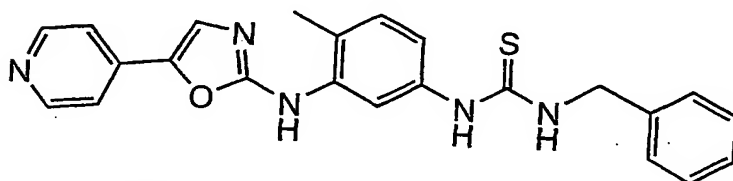
- 024: (2-{2-Methyl-5-[3-(4-trifluoromethyl-phenyl)-ureido]-phenylamino}-oxazol-5-yl)-acetic acid ethyl ester

20



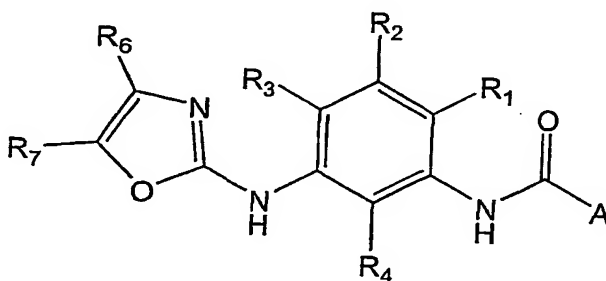
m.p. = 185°C

- 5 025: 1-Benzyl-3-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-thiourea



m.p. = 219°C

- 10 Among the compounds of formula II, the invention is particularly embodied by the compounds wherein R5 = H, Y is an oxygen and Z is an aryl¹ group, corresponding to the N-[3-(Oxazol-2-ylamino)-phenyl]-amide family and the following formula II-2 :



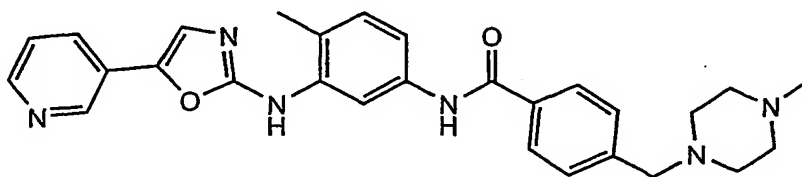
15

FORMULA II-2

- 20 Wherein A is aryl¹ or heteroaryl¹ and wherein R1, R2, R3, R4, R6, R7, aryl¹, heteroaryl¹ have the meaning described on pages as defined in formula I.

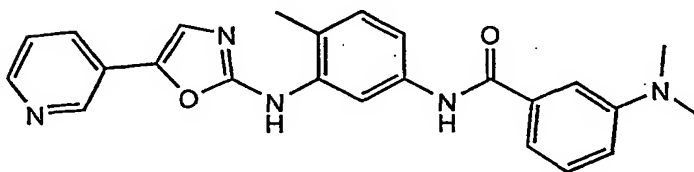
Examples

- 009: 4-(4-Methyl-piperazin-1-ylmethyl)-N-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-benzamide



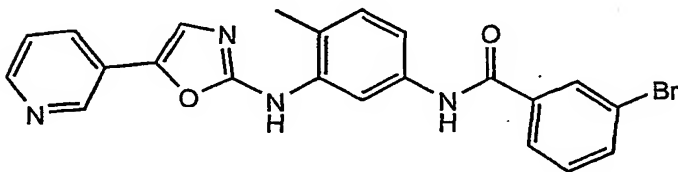
m.p. = 218°C

- 010: 3-Dimethylamino-N-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-benzamide



m.p. = 215°C

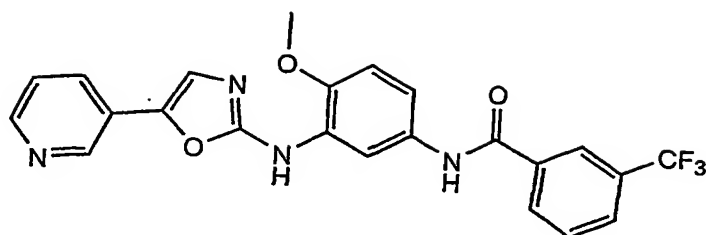
- 011: 3-Bromo-N-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-benzamide



m.p. = 244°C

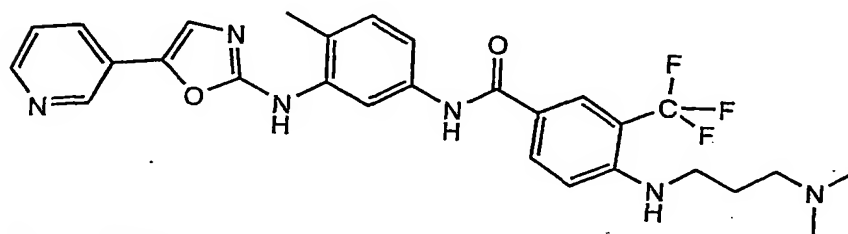
- 026: N-[4-Methoxy-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-trifluoromethyl-benzamide

20



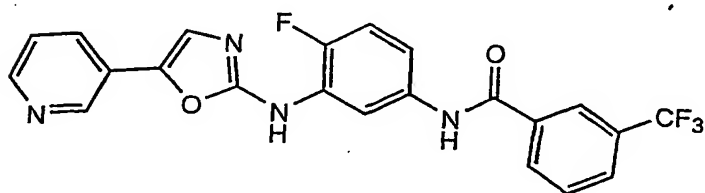
m.p. = 229°C

- 5 027: 4-(3-Dimethylamino-propylamino)-*N*-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-trifluoromethyl-benzamide



10 m.p. = 247°C

- 028: *N*-[4-Fluoro-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-trifluoromethyl-benzamide



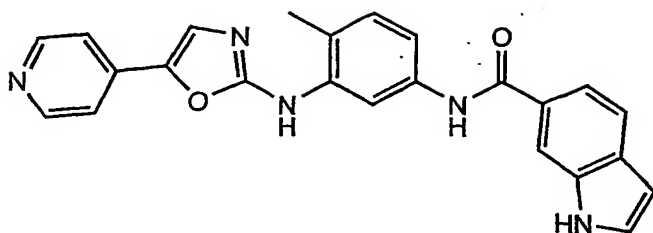
15

m.p. > 265 °C

- 029: 1*H*-Indole-6-carboxylic acid [4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-amide

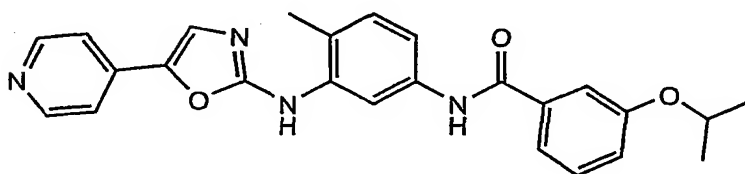
20

21



m.p. = 230-240°C

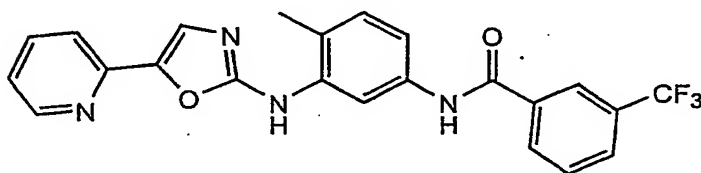
- 5 030: 3-Isopropoxy-*N*-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-benzamide



m.p. = 179-181°C

10

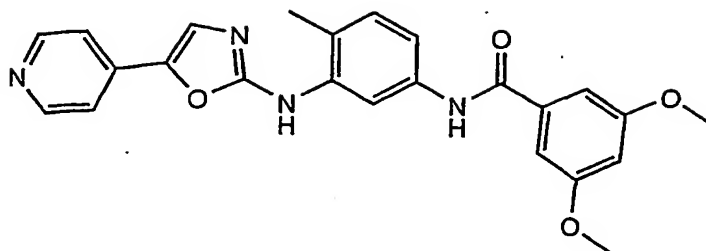
- 031: *N*-[4-Methyl-3-(5-pyridin-2-yl-oxazol-2-ylamino)-phenyl]-3-trifluoromethyl-benzamide



- 15 m.p. = 259°C

- 032: 3,5-Dimethoxy-*N*-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-benzamide

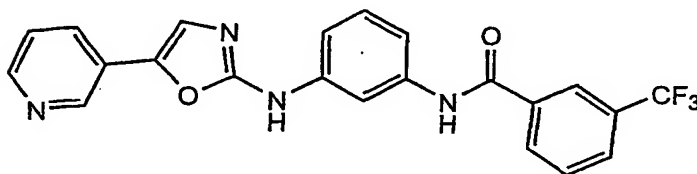
22



m.p. = 209°C

033: *N*-[3-(5-Pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-trifluoromethyl-benzamide

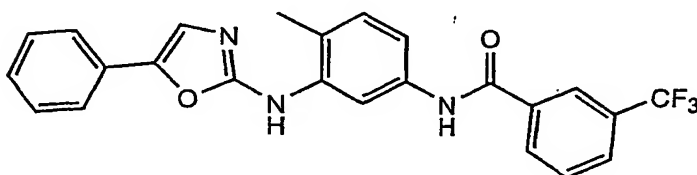
5



m.p. > 265°C

034: *N*-[4-Methyl-3-(5-phenyl-oxazol-2-ylamino)-phenyl]-3-trifluoromethyl-benzamide

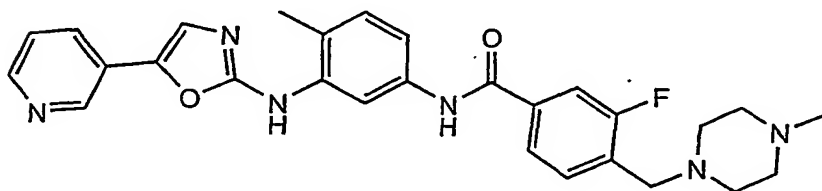
10



m.p. = 212°C

035: 3-Fluoro-4-(4-methyl-piperazin-1-ylmethyl)-*N*-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-benzamide

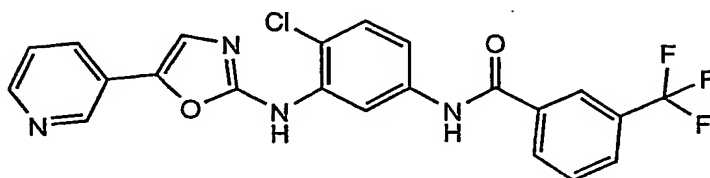
15



20

m.p. = 209°C

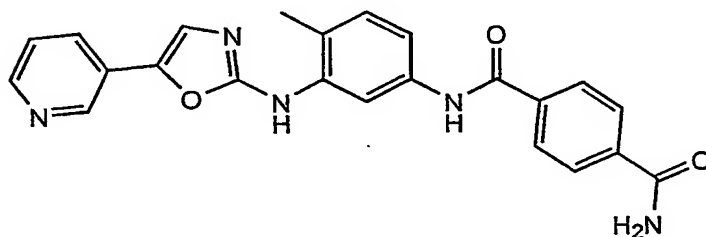
036: *N*-[4-Chloro-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-trifluoromethyl-
5 benzamide



m.p. = 206°C

10

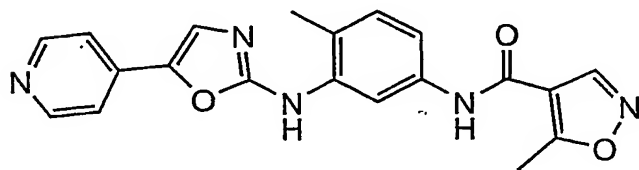
037: *N*-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-terephthalamide



15 m.p. > 265°C

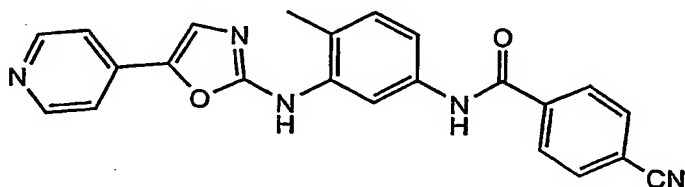
038: 5-Methyl-isoxazole-4-carboxylic acid [4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-amide

20



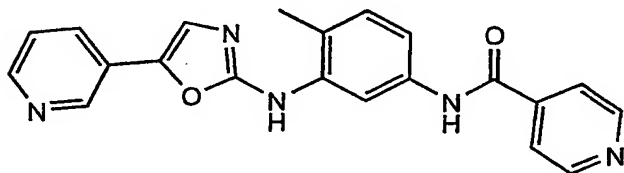
m.p. = 218°C

039: 4-Cyano-*N*-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-benzamide



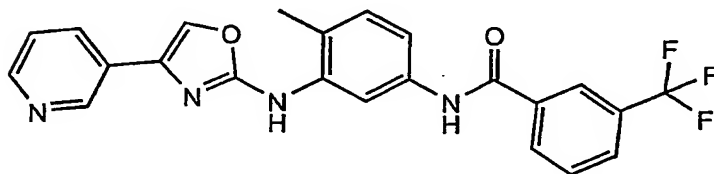
m.p. > 265°C

040: *N*-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-isonicotinamide



m.p. = 249°C

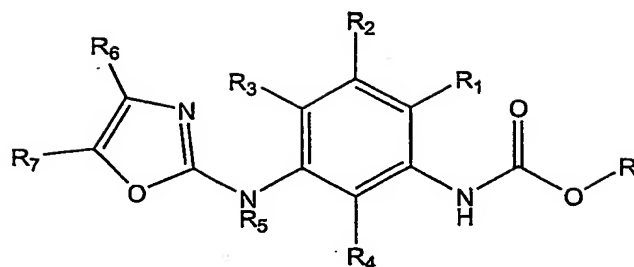
041: *N*-[4-Methyl-3-(4-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-trifluoromethyl-benzamide



m.p. = 245°C

Among the compounds of formula II, the invention is particularly embodied by the compounds wherein Y = O and Z a OR group, corresponding to the family [3-(Oxazol-2-ylamino)-phenyl]-carbamate and the following formula II-3.

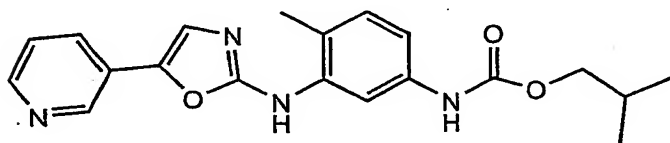
25

**FORMULA II-3**

- 5 wherein R is independently alkyl¹, aryl¹ or heteroaryl¹. R1, R2, R3, R4, R5, R6, and R7 have the meaning described above for formula I.

Examples

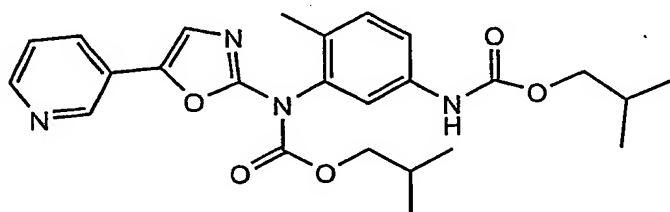
- 10 012: [4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-carbamic acid isobutyl ester



m.p. = 186°C

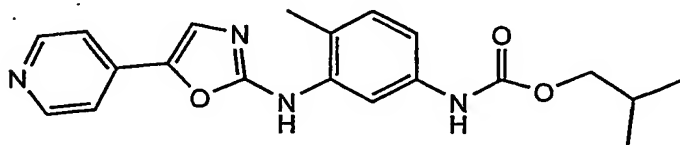
15

- 013: (5-Isobutoxycarbonylamino-2-methyl-phenyl)-(5-pyridin-3-yl-oxazol-2-yl)-carbamic acid isobutyl ester



20 m.p. = 194°C

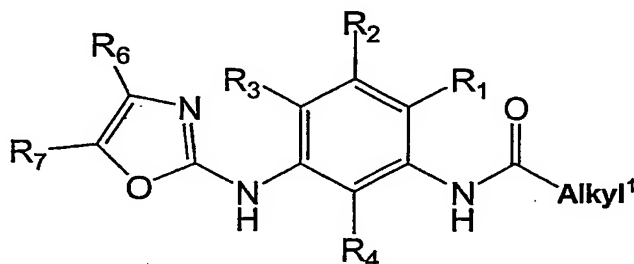
042: [4-Methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-carbamic acid isobutyl ester



5 m.p. = 170°C

Among the compounds of formula II, the invention is particularly embodied by the compounds wherein R₅ = H, Y is an oxygen and Z an alkyl¹ group, corresponding to the family [3-(Oxazol-2-ylamino)-phenyl]-acetamide and the following formula II-4.

10



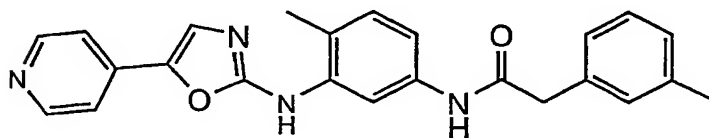
FORMULA II-4

15

R₁, R₂, R₃, R₄, R₆, R₇ and alkyl¹ have the meaning as defined above.

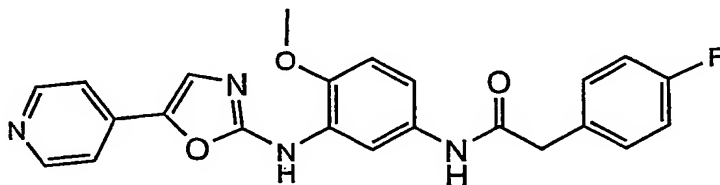
Examples

20 043: *N*-[4-Methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-2-*m*-tolyl-acetamide



m.p. = 212°C

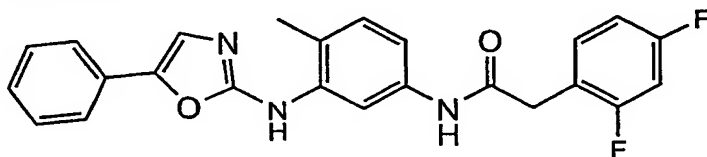
044: 2-(4-Fluoro-phenyl)-*N*-[4-methoxy-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-acetamide



m.p. = 222°C

5

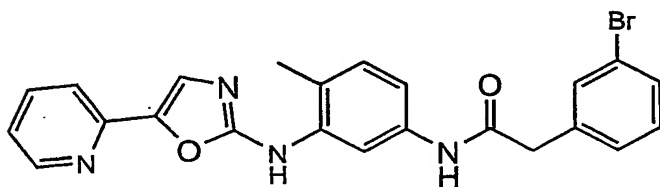
045: 2-(2,4-Difluoro-phenyl)-*N*-[4-methyl-3-(5-phenyl-oxazol-2-ylamino)-phenyl]-acetamide



m.p. = 230°C

10

046: 2-(3-Bromo-phenyl)-*N*-[4-methyl-3-(5-pyridin-2-yl-oxazol-2-ylamino)-phenyl]-acetamide



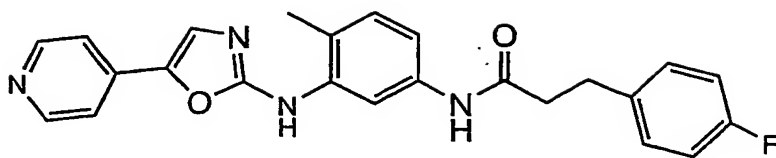
15

m.p. = 211°C

047: 3-(4-Fluoro-phenyl)-*N*-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-propionamide

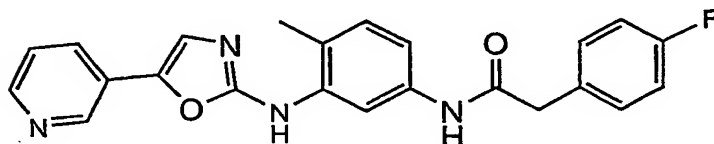
20

28



m.p. = 224°C

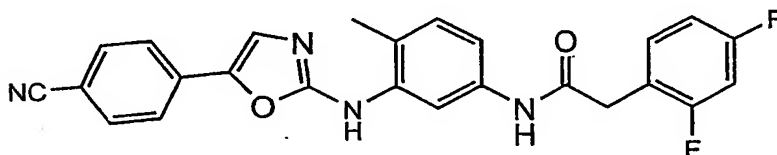
- 048: 2-(4-Fluoro-phenyl)-N-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-
5. acetamide



m.p. = 231-233°C

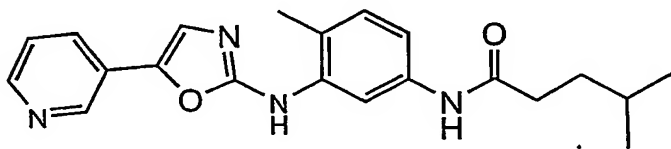
10

- 049: N-{3-[5-(4-Cyano-phenyl)-oxazol-2-ylamino]-4-methyl-phenyl}-2-(2,4-difluoro-
phenyl)-acetamide



15 m.p. = 240°C

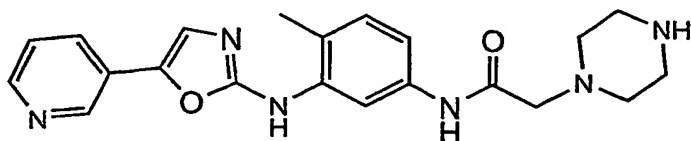
- 050: 4-Methyl-pentanoic acid [4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-
amide



20

m.p. = 211°C

051: *N*-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-2-piperazin-1-yl-acetamide

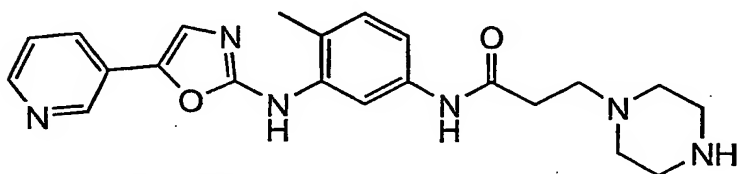


5

m.p. = 176-178°C

052: *N*-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-piperazin-1-yl-propionamide

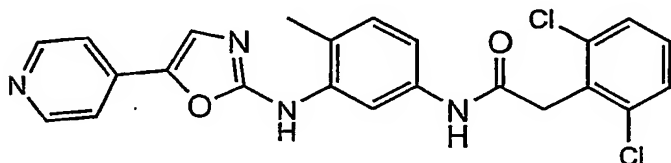
10



m.p. = 118-120°C

053: 2-(2,6-Dichloro-phenyl)-*N*-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-acetamide

15

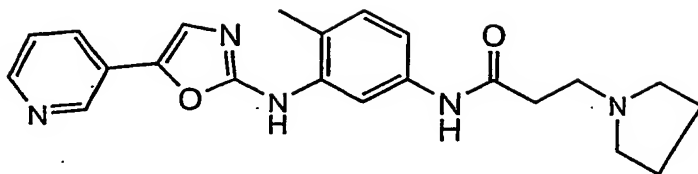


20 m.p. > 265°C

054: *N*-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-3-pyrrolidin-1-yl-propionamide

25

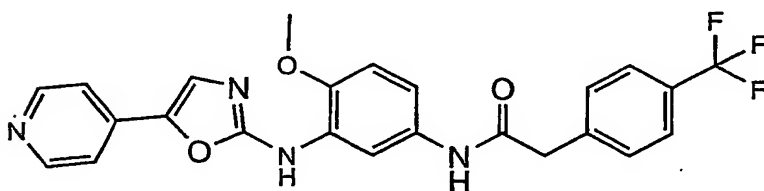
30



m.p. = 186°C

5

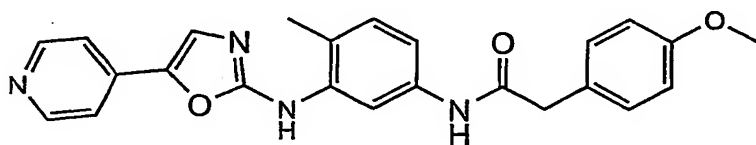
055: *N*-[4-Methoxy-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-2-(4-trifluoromethyl-phenyl)-acetamide



m.p. = 225°C

15

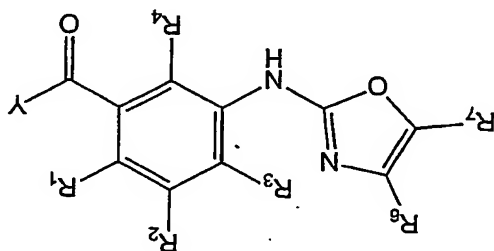
056: 2-(4-Methoxy-phenyl)-*N*-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-acetamide



m.p. = 187°C

25 Among the compounds of formula I, the invention is particularly embodied by the compounds of the following formula III :

31



FORMULA III

5

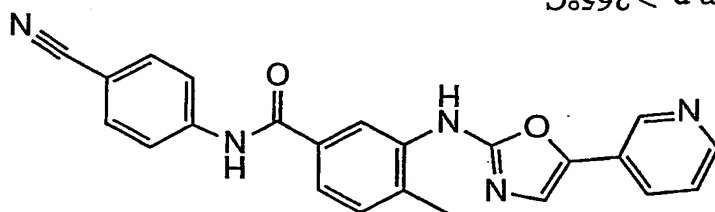
Wherein Y is selected from NR_aR_b, NHNR_aR_b, alkyl¹, aryl¹, or O-R_a wherein R_a and R_b are independently chosen from H or alkyl¹ or aryl¹ or heteroaryl¹, optionally substituted by a pendant basic nitrogen functionality. R₁, R₂, R₃, R₄, R₆, and R₇ have the meaning described above for formula I.

10

Examples:

15

058: *N*-(4-Cyano-phenyl)-4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-benzamide

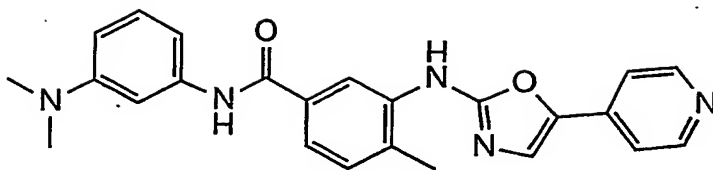


20

m.p. > 265°C

059: *N*-(3-Dimethylamino-phenyl)-4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-benzamide

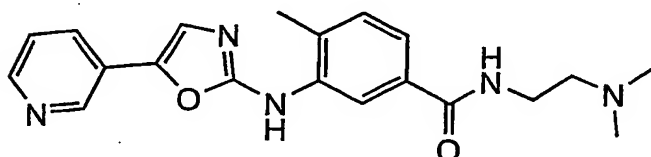
25



m.p. = 230°C

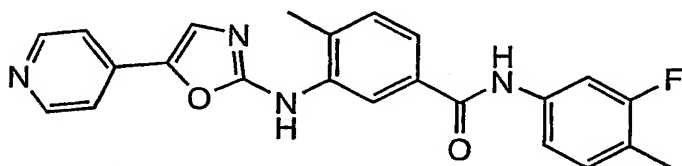
060: *N*-(2-Dimethylamino-ethyl)-4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-benzamide

5



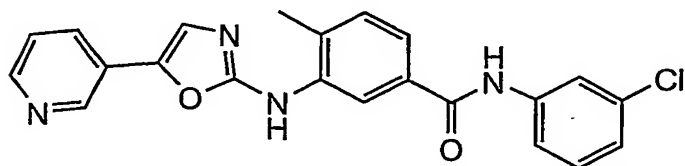
¹H NMR (CDCl₃, 300 MHz) δ = 2.21 (s, 6H) ; 2.33 (s, 3H) ; 2.48 (t, J = 5.9 Hz, 2H) ;
10 3.47 (q, J = 5.6 Hz, 2H) ; 6.82 (s, 1H) ; 6.93 (s, 1H) ; 7.20 (m, 2H) ; 7.25 (m, 1H) ;
7.40 (dd, J = 7.6-1.5 Hz, 1H) ; 7.74 (dt, J = 8.0-1.8 Hz, 1H) ; 8.41 (dd, J = 6.9-1.3 Hz,
2H) ; 8.76 (d, J = 1.8, 1H).

15 061: *N*-(3-Fluoro-4-methyl-phenyl)-4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-benzamide



20 m.p. = 203°C

062: *N*-(3-Chloro-phenyl)-4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-benzamide



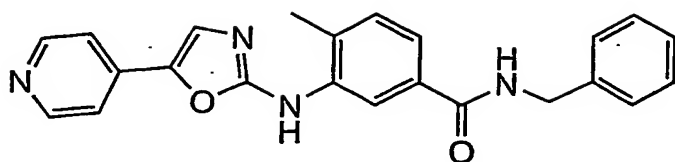
25

m.p. = 247°C

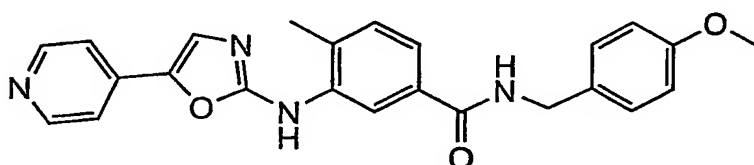
063: *N*-Benzyl-4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-benzamide

30

33

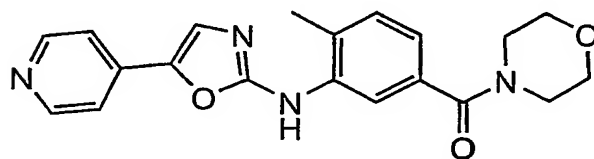


m.p. = 212°C

5 064: *N*-(4-Methoxy-benzyl)-4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-benzamide

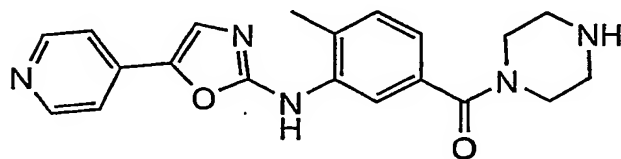
10 m.p. = 212-214°C

065: [4-Methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-morpholin-4-yl-methanone



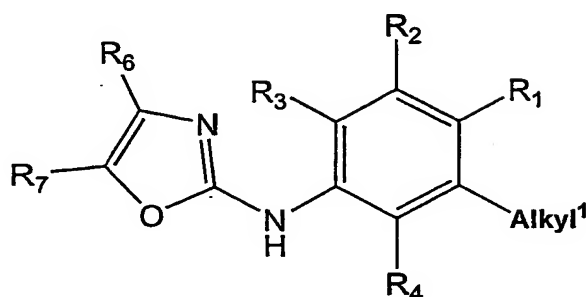
20 m.p. = 155°C

066: [4-Methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-piperazin-1-yl-methanone



25 m.p. = 171°C

Among the compounds of formula I, the invention is particularly embodied by the compounds of the following formula IV :

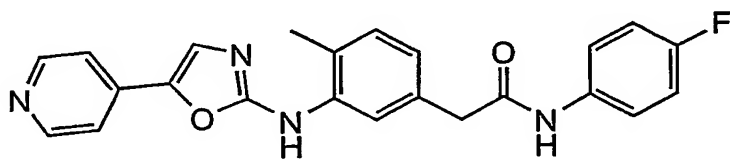


FORMULA IV

Wherein alkyl¹, R₁, R₂, R₃, R₄, R₆, and R₇ have the meaning as defined for formula I above.

Examples:

067: *N*-(4-Fluoro-phenyl)-2-[4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-phenyl]-acetamide



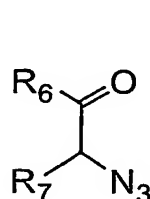
m.p. = 220-222°C

Among the compounds as described above of formula I, I-2, I-3, II, II-2, II-3, II-4, III and IV, the invention contemplates more particularly the groups wherein R₆ is hydrogen and R₇ is pyridyl; which pyridyl may additionally bear any combination, at any one ring position, of one, two or three or four substituents such as

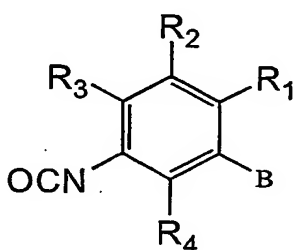
- halogen (selected from F, Cl, Br or I);
- an alkyl¹ group;

- an aryl¹ group;
- trifluoromethyl, O-alkyl¹, carboxyl, cyano, nitro, formyl, hydroxy, NH-alkyl¹, N(alkyl¹)(alkyl¹), and amino, the latter nitrogen substituents optionally in the form of a basic nitrogen functionality;
- 5 - NHCO-R or NHCOO-R or NHCONH-R or NHSO₂-R or NHSO₂NH-R or CO-R or COO-R or CONH-R or SO₂-R or SO₂NH-R wherein R corresponds to hydrogen, alkyl¹ or aryl¹ group.

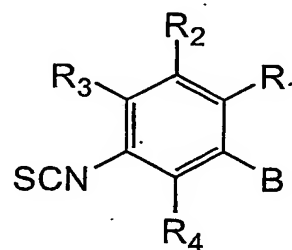
10 In a second embodiment, the invention is directed to a process for manufacturing a compound of formula I depicted above. This entails the condensation of an azide of general formula 10 with an isocyanate of the type 11 or an isothiocyanate of the type 12.



10



11



12

15

Group B in formula 11 and 12 corresponds to group X as described in formula I as well as NO₂, CO₂, and CH₃. R₁, R₂, R₃, R₄, R₆ and R₇ have the meaning described above for formula I.

20

The reaction of 10 either with 11 or 12 in a solvent such as methylene chloride or dioxane in the presence of triphenylphosphine, leads to an oxazole-type product of formula 13.

The following examples are intended to illustrate the present invention.

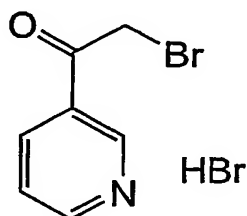
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Example of Compound synthesis

General: All chemicals used were commercial reagent grade products. Solvents were of anhydrous commercial grade and were used without further purification.

- 5 Dichloromethane and dioxane were freshly distilled under a stream of argon before use. The progress of the reactions was monitored by thin layer chromatography using precoated silica gel 60F 254, Merck TLC plates, which were visualized under UV light. Multiplicities in ^1H NMR spectra are indicated as singlet (s), broad singlet (br s), doublet (d), triplet (t), quadruplet (q), and multiplet (m) and the NMR spectrum
10 were realized on a 300 MHz Bruker spectrometer.

Preparation of 3-Bromoacetylpyridine, HBr salt



15

- Bromine (24 g, 150 mmol) in 4 mL of 45% HBr was added dropwise under vigorous stirring to a solution at 70°C of 3-acetyl-pyridine (18 g, 148 mmol) in acetic acid containing 45% of HBr (165 mL). The vigorously stirred mixture was keep at 70°C for 3h. The mixture was cooled and the precipitate collected by filtration and washed
20 with petroleum ether/methanol (1/1, 100 mL) to give 35.8 g of a white crystals (85%).
m.p. = 189°C

^1H NMR (DMSO-*d*₆) δ = 5.09 (s, 2H, CH₂Br) ; 7.78-7.96 (m, 1H, pyridyl-H) ; 8.42-8.70 (m, 1H, pyridyl-H); 8.79-8.99 (m, 1H, pyridyl-H) ; 9.29 (m, 1H, pyridyl-H) ; 12.77 (br s, 1H, HBr)

25

Preparation of 3-Azidoacetylpyridine

37

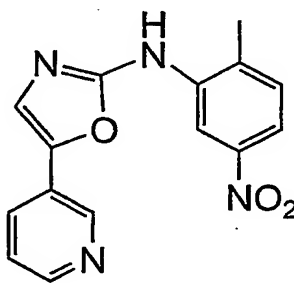


To a solution of 3-bromoacetylpyridine hydrobromide (5 g, 17.8 mmol) in 20 mL of water was added sodium azide (1.16 g, 17.8 mmol) and the contents stirred at room temperature for 2h. The reaction mixture was treated with saturated aqueous NaHCO₃ until neutrality, extracted with ethyl acetate (3x30mL) and the combined organic phases were dried over MgSO₄. After solvent removal the crude residue was silica gel column chromatographed (dichloromethane/ethanol / 98/2). 3-Azidoacetyl-pyridine was obtained as yellow solid (2.17 g, 71%).

m.p. = 69-71°C

¹H NMR (CDCl₃) δ = 4.50 (s, 2H, CH₂N₃); 7.38 (dd, *J* = 7.9-4.9, 1H, pyridyl-H); 8.11 (d, *J*=7.9, 1H, pyridyl-H); 8.71 (d, *J* = 7.9, 1H, pyridyl-H); 8.99 (s, 1H, pyridyl-H).

Preparation of (2-Methyl-5-nitro-phenyl)-(5-pyridin-3-yl-oxazol-2-yl)-amine



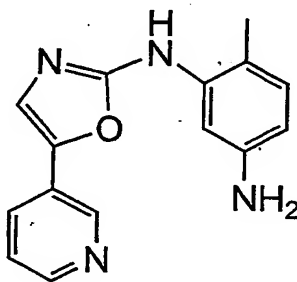
To a solution of 3-azidoacetylpyridine (800 mg, 4.94 mmol) in dioxane 10 mL was added 2-methyl-5-nitrophenyl isocyanate (880 mg, 4.94 mmol) (commercially available), and triphenylphosphine (1.29 g, 4.94 mmol). The reaction mixture was placed in an oil bath preheated to 100°C and stirred for 30 min. After evaporation of the solvent under reduced pressure the residue was partitioned between 4N HCl (20

mL) and dichloromethane (20 mL). The aqueous layer was neutralized with 15% NaOH and extracted with dichloromethane (3x20 mL). The combined organic layers were washed with brine (20 mL), dried over anhydrous MgSO_4 , concentrated. The residue was silica gel column chromatographed (dichloromethane/ethanol : 97/3) to give the title compound as yellow micro crystals (1.14 g, 78%).

m.p. = 252°C

^1H NMR ($\text{DMSO}-d_6$) δ = 2.46 (s, 3H, ArCH_3) ; 7.47-7.50 (m, 2H) ; 7.71 (s, 1H) ; 7.81 (dd, J = 8.2-2.4, 1H) ; 7.98 (d, J = 7.6, 1H) ; 8.48 (d, J = 4.6, 1H) ; 8.88 (br s, 1H) ; 9.07 (br s, 1H) ; 9.62 (s, 1H, NH).

Preparation of (2-Methyl-5-amino-phenyl)-(5-pyridin-3-yl-oxazol-2-yl)-amine

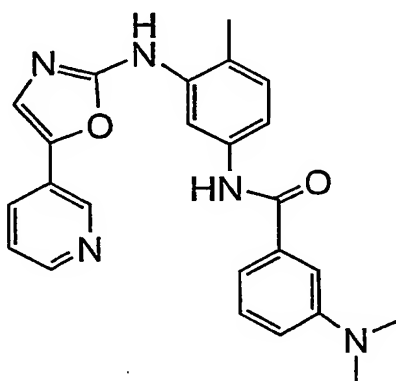


To a solution of (2-methyl-5-nitro-phenyl)-(5-pyridin-3-yl-oxazol-2-yl)-amine (600 mg, 2.02 mmol) in ethanol (20 mL) was added tin(II) chloride dihydrate (2.50 g, 10 mmol). The reaction mixture was heated under reflux for 6h. The mixture was then concentrated, saturated aqueous NaHCO_3 was added and the resultant suspension was extracted with ethyl acetate (3x20 mL). The combined organic layers were washed with brine (20 mL), dried over anhydrous MgSO_4 and concentrated. The residue was silica gel column chromatographed (dichloromethane/ethanol : 97/3). 350 mg (65%) of (2-methyl-5-amino-phenyl)-(5-pyridin-3-yl-oxazol-2-yl)-amine was obtained as pale yellow powder.

m.p. = 166°C

^1H NMR (CDCl_3) δ = 2.17 (s, 3H, ArCH_3) ; 6.28-6.31 (m, 1H) ; 6.63 (br s, 1H) ; 6.90 (d, J = 8.1, 1H) ; 7.18 (s, 1H) ; 7.24 (dd, J = 8.0-5.0, 1H) ; 7.44 (d, J = 2.1, 1H) ; 7.72-7.75 (m, 1H) ; 8.41-8.43 (m, 1H) ; 8.76 (br s, 1H).

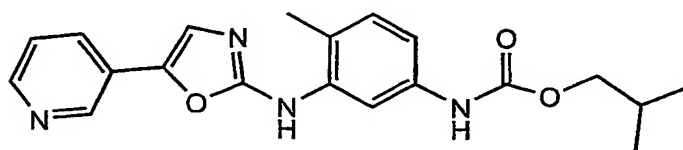
5 Preparation of 3-Dimethylamino-*N*-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-benzamide



- 10 To a solution of (2-methyl-5-amino-phenyl)-(5-pyridin-3-yl-oxazol-2-yl)-amine (120 mg, 0.451 mmol) and 3-dimethylaminobenzoic acid (86 mg, 0.521 mmol) in DMF (6 mL) was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (136 mg, 0.824 mmol), 1-hydroxybenzotriazole (84 mg, 0.622 mmol) and triethylamine (0.98 mL, 0.710 mmol). The mixture was stirred at room temperature for 4h. After
- 15 removal of the solvent, the residue was treated with saturated aqueous NaHCO_3 (20 mL) and extracted with dichloromethane (3×10 mL). The combined organic layers were washed with brine (20 mL), dried over MgSO_4 and concentrated. 3-dimethylamino-*N*-[4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-benzamide was obtained after silica gel column chromatography (dichloromethane/ethanol : 98/2)
- 20 (113 mg, 58%) as yellow solid.
- m.p. = 228°C

^1H NMR ($\text{DMSO}-d^6$) δ = 2.27 (s, 3H, ArCH_3) ; 2.96 (s, 6H, $2\times\text{NCH}_3$) ; 6.90 (d, J = 8.0, 1H) ; 7.14-7.33 (m, 4H) ; 7.40-7.46 (m, 2H) ; 7.57 (s, 1H) ; 7.95 (d, J = 8.0, 1H) ; 8.23 (br s, 1H) ; 8.43 (d, J = 4.7, 1H) ; 8.83 (br s, 1H) ; 9.39 (s, 1H) ; 10.04 (s, 1H).

- 5 Preparation of [4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-carbamic acid isobutyl ester



10

The (2-methyl-5-amino-phenyl)-(5-pyridin-3-yl-oxazol-2-yl)-amine (170 mg, 0.639 mmol) was dissolved in dry THF (7 mL) under argon atmosphere and Chloroformate isobutyl (1.1eq.) was added dropwise at 0°C . The reaction mixture was brought to room temperature and stirred for 3h. Evaporation to dryness gave a solid residue which was purified by alumina gel column chromatography (dichloromethane/ethanol : 98/2) to give white solid (62%).

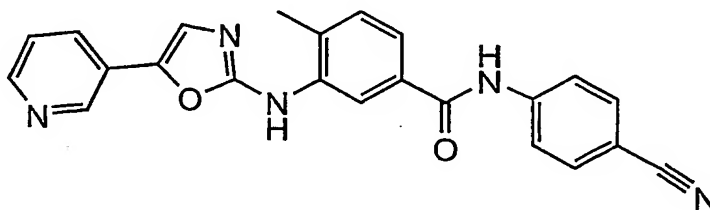
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m.p. = 186°C

^1H NMR ($\text{DMSO}-d^6$) δ = 0.93 (d, J = 6.2, 6H) ; 1.92 (m, 1H) ; 2.21 (s, 3H) ; 3.86 (d, J = 6.2, 2H) ; 7.10 (s, 2H) ; 7.44 (br s, 1H) ; 7.57 (s, 1H) ; 7.95 (s, 1H) ; 8.44 (s, 2H) ; 8.85 (s, 1H) ; 9.35 (s, 1H) ; 9.57 (s, 1H).

20

Preparation of *N*-(4-Cyano-phenyl)-4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-benzamide



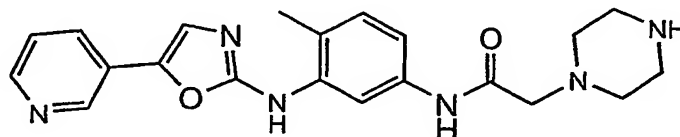
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A 2M solution of trimethyl aluminium in hexanes (4 mL) was added dropwise to a cold (0° C) solution of 4-amino-benzonitrile (236 mg, 2 mmol) in anhydrous dichloromethane (20 mL) under argon atmosphere. The mixture was warmed to room temperature and stirred at room temperature for 3h. A solution of 4-methyl-3-(5-pyridin-4-yl-oxazol-2-ylamino)-benzoic acid methyl ester (620 mg, 2 mmol) in anhydrous dichloromethane (5 mL) and added slowly, and the resulting mixture was heated at reflux for 12h. The mixture was cooled to 0°C and quenched by dropwise addition of a 4N aqueous sodium hydroxide solution (10 mL). The mixture was extracted with dichloromethane (3×30 mL). The combined organic layers were washed with brine (3×40 mL) and dried over anhydrous MgSO₄. *N*-(4-cyano-phenyl)-4-methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-benzamide is obtained in 78% after purification by silica gel column chromatography (dichloromethane/ethanol : 98/2).

m.p. = 203°C

¹H NMR (DMSO-d₆) δ = 2.39 (s, 3H) ; 7.39 (d, *J* = 8.0, 1H) ; 7.46 (m, 1H) ; 7.61 (s, 1H) ; 7.64 (s, 1H) ; 7.82 (d, *J* = 8.6, 2H) ; 7.95 (d, *J* = 8.1, 1H) ; 7.99 (d, *J* = 8.6, 2H) ; 8.45 (s, 1H) ; 8.48 (s, 1H) ; 8.84 (s, 1H) ; 9.61 (s, 1H) ; 10.59 (s, 1H).

preparation of *N*-[4-Methyl-3-(5-pyridin-3-yl-oxazol-2-ylamino)-phenyl]-2-piperazin-1-yl-acetamide



The (2-methyl-5-amino-phenyl)-(5-pyridin-3-yl-oxazol-2-yl)-amine (500 mg, 1.88 mmol) was dissolved in acetone (200 mL) under an argon atmosphere. K₂CO₃ (1.2 eq) was added and the suspension was cooled to 0°C. The chloro-acetyl chloride (1.2 eq) was added dropwise and the mixture was brought to room temperature under stirring for 10h. The resulting yellow suspension was filtered and K₂CO₃ was washed with methanol. The resulting organic solution was evaporated to dryness. The expected

product was taken up from Et₂O to give a yellow powder (90%). To this chloroderivative (200 mg, 0.583 mmol) dissolved in absolute ethanol was added NaI (0.7 eq). The reaction mixture was heated to reflux and piperazine (12 eq) was added. The mixture was stirred for 10h at reflux temperature. The solution was then diluted
5 with dichloromethane and washed with an aqueous saturated solution of NaHCO₃. The aqueous layer was extracted with dichloromethane (3 x 25 ml). The combined extracts were dried over MgSO₄, filtered and concentrated under reduced pressure to dryness. A column chromatography on alumina (dichloromethane/ethanol : 98/2) provided the expected product as a beige solid (70%).

10 m.p. = 176-178°C

¹H NMR (CDCl₃) δ = 3.73 (s, 3H) ; 4.15 (br s, 4H) ; 4.50 (br s, 4H) ; 4.68 (s, 2H) ; 8.61 (d, *J* = 8.3, 1H) ; 8.78 (d, *J* = 7.2, 1H) ; 8.93 (m, 1H) ; 9.06 (s, 1H) ; 9.43 (d, *J* = 7.8, 1H) ; 9.58 (s, 1H) ; 9.93 (dd, *J* = 4.6-1.2, 1H) ; 10.33 (s, 1H) ; 10.87 (s, 1H) ; 11.22 (s, 1H).

15 In a third embodiment, the invention relates to a pharmaceutical composition comprising a compound as depicted above.

Such medicament can take the form of a pharmaceutical composition adapted for oral
20 administration, which can be formulated using pharmaceutically acceptable carriers well known in the art in suitable dosages. Such carriers enable the pharmaceutical compositions to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions, and the like, for ingestion by the patient. In addition to the active ingredients, these pharmaceutical compositions may contain suitable
25 pharmaceutically-acceptable carriers comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations which can be used pharmaceutically. Further details on techniques for formulation and administration may be found in the latest edition of Remington's Pharmaceutical Sciences (Maack Publishing Co., Easton, Pa.).

The composition of the invention can also take the form of a pharmaceutical or cosmetic composition for topical administration.

Such compositions may be presented in the form of a gel, paste, ointment, cream,
5 lotion, liquid suspension aqueous, aqueous-alcoholic or, oily solutions, or dispersions
of the lotion or serum type, or anhydrous or lipophilic gels, or emulsions of liquid or
semi-solid consistency of the milk type, obtained by dispersing a fatty phase in an
aqueous phase or vice versa, or of suspensions or emulsions of soft, semi-solid
consistency of the cream or gel type, or alternatively of microemulsions, of
10 microcapsules, of microparticles or of vesicular dispersions to the ionic and/or
nonionic type. These compositions are prepared according to standard methods.

The composition according to the invention comprises any ingredient commonly used
in dermatology and cosmetic. It may comprise at least one ingredient selected from
15 hydrophilic or lipophilic gelling agents, hydrophilic or lipophilic active agents,
preservatives, emollients, viscosity enhancing polymers, humectants, surfactants,
preservatives, antioxidants, solvents, and fillers, antioxidants, solvents, perfumes,
fillers, screening agents, bactericides, odor absorbers and coloring matter.

20 As oils which can be used in the invention, mineral oils (liquid paraffin), vegetable
oils (liquid fraction of shea butter, sunflower oil), animal oils, synthetic oils, silicone
oils (cyclomethicone) and fluorinated oils may be mentioned. Fatty alcohols, fatty
acids (stearic acid) and waxes (paraffin, carnauba, beeswax) may also be used as fatty
substances.

25

As emulsifiers which can be used in the invention, glycerol stearate, polysorbate 60
and the PEG-6/PEG-32/glycol stearate mixture are contemplated.

As hydrophilic gelling agents, carboxyvinyl polymers (carbomer), acrylic copolymers
such as acrylate/alkylacrylate copolymers, polyacrylamides, polysaccharides such as
30 hydroxypropylcellulose, clays and natural gums may be mentioned, and as lipophilic
gelling agents, modified clays such as bentones, metal salts of fatty acids such as

aluminum stearates and hydrophobic silica, or alternatively ethylcellulose and polyethylene may be mentioned.

As hydrophilic active agents, proteins or protein hydrolysates, amino acids, polyols, urea, allantoin, sugars and sugar derivatives, vitamins, starch and plant extracts, in particular those of Aloe vera may be used.

As lipophilic active agents, retinol (vitamin A) and its derivatives, tocopherol (vitamin E) and its derivatives, essential fatty acids, ceramides and essential oils may be used. These agents add extra moisturizing or skin softening features when utilized.

In addition, a surfactant can be included in the composition so as to provide deeper penetration of the compound capable of depleting mast cells, such as a tyrosine kinase inhibitor, preferably a c-kit and/or a bcr-abl inhibitor.

Among the contemplated ingredients, the invention embraces penetration enhancing agents selected for example from the group consisting of mineral oil, water, ethanol, triacetin, glycerin and propylene glycol; cohesion agents selected for example from the group consisting of polyisobutylene, polyvinyl acetate and polyvinyl alcohol, and thickening agents.

Chemical methods of enhancing topical absorption of drugs are well known in the art. For example, compounds with penetration enhancing properties include sodium lauryl sulfate (Dugard, P. H. and Sheuplein, R. J., "Effects of Ionic Surfactants on the Permeability of Human Epidermis: An Electrometric Study," J. Invest. Dermatol., V.60, pp. 263-69, 1973), lauryl amine oxide (Johnson et. al., US 4,411,893), azone (Rajadhyaksha, US 4,405,616 and 3,989,816) and decylmethyl sulfoxide (Sekura, D. L. and Scala, J., "The Percutaneous Absorption of Alkylmethyl Sulfides," Pharmacology of the Skin, Advances In Biology of Skin, (Appleton-Century Craft) V. 12, pp. 257-69, 1972). It has been observed that increasing the polarity of the head group in amphoteric molecules increases their penetration-enhancing properties but at

the expense of increasing their skin irritating properties (Cooper, E. R. and Berner, B., "Interaction of Surfactants with Epidermal Tissues: Physiochemical Aspects," Surfactant Science Series, V. 16, Reiger, M. M. ed. (Marcel Dekker, Inc.) pp. 195-210, 1987).

5

A second class of chemical enhancers are generally referred to as co-solvents. These materials are absorbed topically relatively easily, and, by a variety of mechanisms, achieve permeation enhancement for some drugs. Ethanol (Gale et. al., U.S. Pat. No. 4,615,699 and Campbell et. al., U.S. Pat. Nos. 4,460,372 and 4,379,454), dimethyl
10 sulfoxide (US 3,740,420 and 3,743,727, and US 4,575,515), and glycerine derivatives (US 4,322,433) are a few examples of compounds which have shown an ability to enhance the absorption of various compounds.

The pharmaceutical compositions of the invention can also be intended for
15 administration with aerosolized formulation to target areas of a patient's respiratory tract.

Devices and methodologies for delivering aerosolized bursts of a formulation of a drug is disclosed in US 5,906,202. Formulations are preferably solutions, e.g. aqueous
20 solutions, ethanoic solutions, aqueous/ethanoic solutions, saline solutions, colloidal suspensions and microcrystalline suspensions. For example aerosolized particles comprise the active ingredient mentioned above and a carrier, (e.g., a pharmaceutically active respiratory drug and carrier) which are formed upon forcing the formulation through a nozzle which nozzle is preferably in the form of a flexible
25 porous membrane. The particles have a size which is sufficiently small such that when the particles are formed they remain suspended in the air for a sufficient amount of time such that the patient can inhale the particles into the patient's lungs.

The invention encompasses the systems described in US 5,556,611:

- liquid gas systems (a liquefied gas is used as propellant gas (e.g. low-boiling FCHC
30 or propane, butane) in a pressure container,

- suspension aerosol (the active substance particles are suspended in solid form in the liquid propellant phase),

- pressurized gas system (a compressed gas such as nitrogen, carbon dioxide, dinitrogen monoxide, air is used.

5 Thus, according to the invention the pharmaceutical preparation is made in that the active substance is dissolved or dispersed in a suitable nontoxic medium and said solution or dispersion atomized to an aerosol, i.e. distributed extremely finely in a carrier gas. This is technically possible for example in the form of aerosol propellant gas packs, pump aerosols or other devices known per se for liquid misting and solid
10 atomizing which in particular permit an exact individual dosage.

Therefore, the invention is also directed to aerosol devices comprising the compound as defined above and such a formulation, preferably with metered dose valves.

The pharmaceutical compositions of the invention can also be intended for intranasal
15 administration.

In this regard, pharmaceutically acceptable carriers for administering the compound to the nasal mucosal surfaces will be readily appreciated by the ordinary artisan. These carriers are described in the Remington's Pharmaceutical Sciences" 16th edition, 1980,
20 Ed. By Arthur Osol, the disclosure of which is incorporated herein by reference.

The selection of appropriate carriers depends upon the particular type of administration that is contemplated. For administration via the upper respiratory tract, the composition can be formulated into a solution, e.g., water or isotonic saline, buffered or unbuffered, or as a suspension, for intranasal administration as drops or as
25 a spray. Preferably, such solutions or suspensions are isotonic relative to nasal secretions and of about the same pH, ranging e.g., from about pH 4.0 to about pH 7.4 or, from pH 6.0 to pH 7.0. Buffers should be physiologically compatible and include, simply by way of example, phosphate buffers. For example, a representative nasal
30 decongestant is described as being buffered to a pH of about 6.2 (Remington's, Id. at page 1445). Of course, the ordinary artisan can readily determine a suitable saline

content and pH for an innocuous aqueous carrier for nasal and/or upper respiratory administration.

Common intranasal carriers include nasal gels, creams, pastes or ointments with a viscosity of, e.g., from about 10 to about 3000 cps, or from about 2500 to 6500 cps, or greater, may also be used to provide a more sustained contact with the nasal mucosal surfaces. Such carrier viscous formulations may be based upon, simply by way of example, alkylcelluloses and/or other biocompatible carriers of high viscosity well known to the art (see e.g., Remington's, cited supra. A preferred alkylcellulose is, e.g., methylcellulose in a concentration ranging from about 5 to about 1000 or more mg per 100 ml of carrier. A more preferred concentration of methyl cellulose is, simply by way of example, from about 25 to about mg per 100 ml of carrier.

Other ingredients, such as art known preservatives, colorants, lubricating or viscous mineral or vegetable oils, perfumes, natural or synthetic plant extracts such as aromatic oils, and humectants and viscosity enhancers such as, e.g., glycerol, can also be included to provide additional viscosity, moisture retention and a pleasant texture and odor for the formulation. For nasal administration of solutions or suspensions according to the invention, various devices are available in the art for the generation of drops, droplets and sprays.

A premeasured unit dosage dispenser including a dropper or spray device containing a solution or suspension for delivery as drops or as a spray is prepared containing one or more doses of the drug to be administered and is another object of the invention. The invention also includes a kit containing one or more unit dehydrated doses of the compound, together with any required salts and/or buffer agents, preservatives, colorants and the like, ready for preparation of a solution or suspension by the addition of a suitable amount of water.

Another aspect of the invention is directed to the use of said compound to manufacture a medicament. In other words, the invention embraces a method for

treating a disease related to unregulated c-kit transduction comprising administering an effective amount of a compound as defined above to a mammal in need of such treatment. It also relates to a method for treating a disease related bcr-abl and/or Flt-3 comprising administering an effective amount of a compound as defined above to a
5 mammal in need of such treatment.

More particularly, the invention is aimed at a method for treating a disease selected from autoimmune diseases, allergic diseases, bone loss, cancers such as leukemia and GIST, tumor angiogenesis, inflammatory diseases, inflammatory bowel diseases
10 (IBD), interstitial cystitis, mastocytosis, infections diseases, metabolic disorders, fibrosis, diabetes and CNS disorders comprising administering an effective amount a compound depicted above to a mammal in need of such treatment.

The above described compounds are useful for manufacturing a medicament for the
15 treatment of diseases related to unregulated c-kit transduction, including, but not limited to:

- neoplastic diseases such as mastocytosis, canine mastocytoma, solid tumours, human gastrointestinal stromal tumor ("GIST"), small cell lung cancer, non-small cell lung cancer, acute myelocytic leukemia, acute lymphocytic
20 leukemia, myelodysplastic syndrome, chronic myelogenous leukemia, colorectal carcinomas, gastric carcinomas, gastrointestinal stromal tumors, testicular cancers, glioblastomas, solid tumors and astrocytomas.
- tumor angiogenesis.
- metabolic diseases such as diabetes mellitus and its chronic complications;
25 obesity; type II diabetes; hyperlipidemias and dyslipidemias; atherosclerosis; hypertension; and cardiovascular disease.
- allergic diseases such as asthma, allergic rhinitis, allergic sinusitis, anaphylactic syndrome, urticaria, angioedema, atopic dermatitis, allergic contact dermatitis, erythema nodosum, erythema multiforme, cutaneous
30 necrotizing venulitis and insect bite skin inflammation and blood sucking parasitic infestation.

- interstitial cystitis.
- bone loss (osteoporosis).
- inflammatory diseases such as rheumatoid arthritis, conjunctivitis, rheumatoid spondylitis, osteoarthritis, gouty arthritis and other arthritic conditions as well as inflammatory muscle disorders;
- 5 - autoimmune diseases such as multiple sclerosis, psoriasis, intestine inflammatory disease, ulcerative colitis, Crohn's disease, rheumatoid arthritis and polyarthritis, local and systemic scleroderma, systemic lupus erythematosus, discoid lupus erythematosus, cutaneous lupus, dermatomyositis, polymyositis, Sjogren's syndrome, nodular panarteritis, autoimmune enteropathy, as well as proliferative glomerulonephritis.
- 10 - graft-versus-host disease or graft rejection in any organ transplantation including kidney, pancreas, liver, heart, lung, and bone marrow.
- Other autoimmune diseases embraced by the invention active chronic hepatitis and chronic fatigue syndrome.
- 15 - subepidermal blistering disorders such as pemphigus.
- Vasculitis.
- HIV infection.
- Plasmodium infection.
- 20 - melanocyte dysfunction associated diseases such as hypermelanosis resulting from melanocyte dysfunction and including lentigines, solar and senile lentigo, Dubreuilh melanosis, moles as well as malignant melanomas. In this regard, the invention embraces the use of the compounds defined above to manufacture a medicament or a cosmetic composition for whitening human skin.
- 25 - CNS disorders such as psychiatric disorders, migraine, pain, memory loss and nerve cells degeneracy. More particularly, the method according to the invention is useful for the treatment of the following disorders: Depression including dysthymic disorder, cyclothymic disorder, bipolar depression, severe or "melancholic" depression, atypical depression, refractory depression,
- 30 seasonal depression, anorexia, bulimia, premenstrual syndrome, post-

- menopause syndrome, other syndromes such as mental slowing and loss of concentration, pessimistic worry, agitation, self-deprecation, decreased libido, pain including, acute pain, postoperative pain, chronic pain, nociceptive pain, cancer pain, neuropathic pain, psychogenic pain syndromes, anxiety disorders including anxiety associated with hyperventilation and cardiac arrhythmias, phobic disorders, obsessive-compulsive disorder, posttraumatic stress disorder, acute stress disorder, generalized anxiety disorder, psychiatric emergencies such as panic attacks, including psychosis, delusional disorders, conversion disorders, phobias, mania, delirium, dissociative episodes including dissociative amnesia, dissociative fugue and dissociative identity disorder, depersonalization, catatonia, seizures, severe psychiatric emergencies including suicidal behaviour, self-neglect, violent or aggressive behaviour, trauma, borderline personality, and acute psychosis, schizophrenia including paranoid schizophrenia, disorganized schizophrenia, catatonic schizophrenia, and undifferentiated schizophrenia,
- neurodegenerative diseases including Alzheimer's disease, Parkinson's disease, Huntington's disease, the prion diseases, Motor Neurone Disease (MND), and Amyotrophic Lateral Sclerosis (ALS).
 - substance use disorders as referred herein include but are not limited to drug addiction, drug abuse, drug habituation, drug dependence, withdrawal syndrome and overdose.
 - Cerebral ischemia
 - Fibrosis
 - Duchenne muscular dystrophy
 - fibrodysplasia
 - ACNE
 - as male contraceptive.

Regarding mastocytosis, the invention contemplates the use of the compounds as defined above for treating the different categories which can be classified as follows:

The category I is composed by two sub-categories (IA and IB). Category IA is made by diseases in which mast cell infiltration is strictly localized to the skin. This category represents the most frequent form of the disease and includes : i) urticaria pigmentosa, the most common form of cutaneous mastocytosis, particularly encountered in children, ii) diffuse cutaneous mastocytosis, iii) solitary mastocytoma and iv) some rare subtypes like bullous, erythrodermic and teleangiectatic mastocytosis. These forms are characterized by their excellent prognosis with spontaneous remissions in children and a very indolent course in adults. Long term survival of this form of disease is generally comparable to that of the normal population and the translation into another form of mastocytosis is rare. Category IB is represented by indolent systemic disease (SM) with or without cutaneous involvement. These forms are much more usual in adults than in children. The course of the disease is often indolent, but sometimes signs of aggressive or malignant mastocytosis can occur, leading to progressive impaired organ function.

The category II includes mastocytosis with an associated hematological disorder, such as a myeloproliferative or myelodysplastic syndrome, or acute leukemia. These malignant mastocytosis does not usually involve the skin. The progression of the disease depends generally on the type of associated hematological disorder that conditions the prognosis.

The category III is represented by aggressive systemic mastocytosis in which massive infiltration of multiple organs by abnormal mast cells is common. In patients who pursue this kind of aggressive clinical course, peripheral blood features suggestive of a myeloproliferative disorder are more prominent. The progression of the disease can be very rapid, similar to acute leukemia, or some patients can show a longer survival time.

Finally, the category IV of mastocytosis includes the mast cell leukemia, characterized by the presence of circulating mast cells and mast cell progenitors representing more than 10% of the white blood cells. This entity represents probably the rarest type of leukemia in humans, and has a very poor prognosis, similar to the rapidly progressing variant of malignant mastocytosis. Mast cell leukemia can occur either *de novo* or as the terminal phase of urticaria pigmentosa or systemic mastocytosis.

The invention also contemplates the method as depicted for the treatment of recurrent bacterial infections, resurging infections after asymptomatic periods such as bacterial cystitis. More particularly, the invention can be practiced for treating FimH expressing bacteria infections such as Gram-negative enterobacteria including *E. coli*, *Klebsiella pneumoniae*, *Serratia marcescens*, *Citrobacter freundii* and *Salmonella typhimurium*.

In this method for treating bacterial infection, separate, sequential or concomitant administration of at least one antibiotic selected bacitracin, the cephalosporins, the penicillins, the aminoglycosides, the tetracyclines, the streptomycins and the macrolide antibiotics such as erythromycin; the fluoroquinolones, actinomycin, the sulfonamides and trimethoprim, is of interest.

In one preferred embodiment, the invention is directed to a method for treating neoplastic diseases such as mastocytosis, canine mastocytoma, solid tumours, human gastrointestinal stromal tumor ("GIST"), small cell lung cancer, non-small cell lung cancer, acute myelocytic leukemia, acute lymphocytic leukemia, myelodysplastic syndrome, chronic myelogenous leukemia, colorectal carcinomas, gastric carcinomas, gastrointestinal stromal tumors, testicular cancers, glioblastomas, and astrocytomas comprising administering a compound as defined herein to a human or mammal, especially dogs and cats, in need of such treatment.

In one other preferred embodiment, the invention is directed to a method for treating allergic diseases such as asthma, allergic rhinitis, allergic sinusitis, anaphylactic

syndrome, urticaria, angioedema, atopic dermatitis, allergic contact dermatitis, erythema nodosum, erythema multiforme, cutaneous necrotizing venulitis and insect bite skin inflammation and blood sucking parasitic infestation comprising administering a compound as defined herein to a human or mammal, especially dogs
5 and cats, in need of such treatment.

In still another preferred embodiment, the invention is directed to a method for treating inflammatory diseases such as rheumatoid arthritis, conjunctivitis, rheumatoid spondylitis, osteoarthritis, gouty arthritis and other arthritic conditions comprising
10 administering a compound as defined herein to a human in need of such treatment.

In still another preferred embodiment, the invention is directed to a method for treating autoimmune diseases such as multiple sclerosis, psoriasis, intestine inflammatory disease, ulcerative colitis, Crohn's disease, rheumatoid arthritis and
15 polyarthritis, local and systemic scleroderma, systemic lupus erythematosus, discoid lupus erythematosus, cutaneous lupus, dermatomyositis, polymyositis, Sjogren's syndrome, nodular panarteritis, autoimmune enteropathy, as well as proliferative glomerulonephritis comprising administering a compound as defined herein to a human in need of such treatment.

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In still another preferred embodiment, the invention is directed to a method for treating graft-versus-host disease or graft rejection in any organ transplantation including kidney, pancreas, liver, heart, lung, and bone marrow comprising administering a compound as defined herein to a human in need of such treatment.

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Example : in vitro TK inhibition assays

• Procedures

○ C-Kit WT and mutated C-Kit (JM and Kinase domain 816) assay

Proliferation assays

30 Cells were washed two times in PBS before plating at 5×10^4 cells per well of 96-well plates in triplicate and stimulated either with hematopoietic growth factors (HGF) or

without. After 2 days of culture, 37 Bq (1.78 Tbq/mmol) of [3 H] thymidine (Amersham Life Science, UK) was added for 6 hours. Cells were harvested and filtered through glass fiber filters and [3 H] thymidine incorporation was measured in a scintillation counter.

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For proliferation assay, all drugs were prepared as 20mM stock solutions in DMSO and conserved at -80°C. Fresh dilutions in PBS were made before each experiment. DMSO dissolved drugs were added at the beginning of the culture. Control cultures were done with corresponding DMSO dilutions. Results are represented in percentage by taking the proliferation without inhibitor as 100%.

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Cells

Ba/F3 murine kit and human kit, Ba/F3 mkit Δ 27 (juxtamembrane deletion), and hkitD816V are derived from the murine IL-3 dependent Ba/F3 proB lymphoid cells. The FMA3 and P815 cell lines are mastocytoma cells expressing endogenous mutated forms of Kit, i.e., frame deletion in the murine juxtamembrane coding region of the receptor-codons 573 to 579. The human leukaemic MC line HMC-1 expresses a double point mutation (i.e. mutations JM-V560G and the kinase domain mutation kitD816V), whereas the HMC1 subclone α 155 expresses only the mutation JM-V560G.

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20 Immunoprecipitation assays and western blotting analysis

For each assay, 5.10 \times 10⁶ Ba/F3 cells and Ba/F3-derived cells with various c-kit mutations were lysed and immunoprecipitated as described (Beslu *et al.*, 1996), excepted that cells were stimulated with 250 ng / ml of rmKL. Cell lysates were immunoprecipitated with rabbit immunsera directed against the KIT cytoplasmic domain either with an anti murine KIT (Rottapel *et al.*, 1991) or an anti human KIT (Santa Cruz). Western blot was hybridized either with the 4G10 anti-phosphotyrosine antibody (UBI) or with the appropriate rabbit immunsera anti KIT or with different antibodies (described in antibodies paragraph). The membrane was then incubated either with HRP-conjugated goat anti mouse IgG antibody or with HRP-conjugated

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goat anti rabbit IgG antibody (Immunotech), Proteins of interest were then visualized by incubation with ECL reagent (Amersham).

- **Experimental results**

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The experimental results for various compounds according to the invention using above-described protocols are set forth at Table 1:

Table 1 : in vitro inhibitions of various compounds against c-Kit WT, c-Kit JM Δ 27 and c-Kit D816V.

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Target	IC ₅₀ (μ M)	Compounds
c-Kit WT	IC ₅₀ < 1 μ M	005 ; 006 ; 008 ; 009 ; 019 ; 010 ; 011 ; 023 ; 025 ; 026 ; 027 ; 028 ; 029 ; 032 ; 033 ; 035 ; 036 ; 038 ; 039 ; 040 ; 041 ; 042 ; 043 ; 044 ; 048 ; 050 ; 055 ; 056 ; 057 ; 059 ; 061 ; 062 ;
c-Kit JM Δ 27	IC ₅₀ < 1 μ M	003 ; 004 ; 005 ; 006 ; 007 ; 008 ; 009 ; 010 ; 011 ; 012 ; 019 ; 022 ; 042 ; 050 ; 055 ;
c-Kit D816V	IC ₅₀ \leq 1 μ M	023 ; 025 ; 026 ; 029 ; 032 ; 035 ; 036 ; 038 ; 039 ; 041 ; 042 ; 043 ; 044 ; 055 ; 059 ; 063 ;